In the Claims

Please cancel claim 62. Please amend claims 13-17 and 39-61 as set out below.

13. (Once Amended) A method for the treatment of a hepatitis B virus infection in a human comprising administering an effective amount of β -L-2'-deoxycytidine of the formula:



or pharmaceutically acceptable salt thereof.

14. (Once Amended) A method for the treatment of a hepatitis B virus infection in a human comprising administering an effective amount of β -L-thymidine of the formula:



or pharmaceutically acceptable salt thereof.

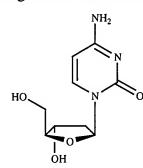




15. (Once Amended) A method for the treatment of a hepatitis B virus infection in a human comprising administering an effective amount of a combination of the following nucleosides:

or pharmaceutically acceptable salt thereof.

16. (Once Amended) A method for the treatment of a hepatitis B virus infection in a human comprising administering an effective amount of a compound of the formula:



or pharmaceutically acceptable salt thereof, in combination or alternation with an effective amount of a compound selected from the group consisting of β -L-2-hydroxymethyl-5-(cytosin-1-yl)-1,3-oxathiolane (3TC), *cis*-2-hydroxymethyl-5-(5-fluorocytosin-1-yl)-1,3-oxathiolane (FTC), β -L-2'-fluoro-5-methyl-arabinofuranosyl-uridine (L-FMAU), β -D-2,6-diaminopurine dioxolane (DAPD), famciclovir, penciclovir, 2-amino-1,9-dihydro-9-[4-hydroxy-3-(hydroxymethyl)-2-methylenecyclopentyl]-6H-purin-6-one (entecavir, BMS-200475), 9-[2-(phosphono-methoxy)ethyl]adenine (PMEA, adefovir, dipivoxil); lobucavir, ganciclovir and ribavirin.





17. (Once Amended) A method for the treatment of a hepatitis B virus infection in a human comprising administering an effective amount of a compound of the formula:

or pharmaceutically acceptable salt thereof, in combination or alternation with an effective amount of a compound selected from the group consisting of β-L-2-hydroxymethyl-5-(cytosin-1-yl)-1,3-oxathiolane (3TC), *cis*-2-hydroxymethyl-5-(5-fluorocytosin-1-yl)-1,3-oxathiolane (FTC), β-L-2'-fluoro-5-methyl-arabinofuranosyl-uridine (L-FMAU), β-D-2,6-diaminopurine dioxolane (DAPD), famciclovir, penciclovir, 2-amino-1,9-dihydro-9-[4-hydroxy-3-(hydroxymethyl)-2-methylenecyclopentyl]-6H-purin-6-one (entecavir, BMS-200475), 9-[2-(phosphono-methoxy)ethyl]adenine (PMEA, adefovir, dipivoxil); lobucavir, ganciclovir and ribavirin.

39. (Once Amended) The method of claim 37 or 38, wherein the dosage unit is a tablet or capsule.

(Once Amended) A method for the treatment of a hepatitis B virus infection in a human comprising administering an effective amount of a compound of the formula:





or pharmaceutically acceptable salt thereof, in combination or alternation with an effective amount of β -L-2-hydroxymethyl-5-(cytosin-1-yl)-1,3-oxathiolane (3TC), or pharmaceutically acceptable salt thereof.

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(Once Amended) A method for the treatment of a hepatitis B virus infection in a human comprising administering an effective amount of a compound of the formula:

or pharmaceutically acceptable salt thereof, in combination or alternation with an effective amount of *cis*-2-hydroxymethyl-5-(5-fluorocytosin-1-yl)-1,3-oxathiolane (FTC), or pharmaceutically acceptable salt thereof.

(Once Amended) A method for the treatment of a hepatitis B virus infection in a human comprising administering an effective amount of a compound of the formula:

or pharmaceutically acceptable salt thereof, in combination or alternation with an effective amount of β -L-2'-fluoro-5-methyl-arabinofuranosyl-uridine (L-FMAU), or pharmaceutically acceptable salt thereof.





14/3.

(Once Amended) A method for the treatment of a hepatitis B virus infection in a human comprising administering an effective amount of a compound of the formula:

or pharmaceutically acceptable salt thereof, in combination or alternation with an effective amount of β -D-2,6-diaminopurine dioxolane (DAPD), or pharmaceutically acceptable salt thereof.

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(Once Amended) A method for the treatment of a hepatitis B virus infection in a human comprising administering an effective amount of a compound of the formula:

or pharmaceutically acceptable salt thereof, in combination or alternation with an effective amount of famciclovir, or pharmaceutically acceptable salt thereof.

10 48.

(Once Amended) A method for the treatment of a hepatitis B virus infection in a human comprising administering an effective amount of a compound of the formula:





or pharmaceutically acceptable salt thereof, in combination or alternation with an effective amount of penciclovir, or pharmaceutically acceptable salt thereof.

36.

(Once Amended) A method for the treatment of a hepatitis B virus infection in a human comprising administering an effective amount of a compound of the formula:

or pharmaceutically acceptable salt thereof, in combination or alternation with an effective amount of 2-amino-1,9-dihydro-9-[4-hydroxy-3-(hydroxymethyl)-2-methylene-cyclopentyl]-6H-purin-6-one (entecavir, BMS-200475), or pharmaceutically acceptable salt thereof.

(Once Amended) A method for the treatment of a hepatitis B virus infection in a human comprising administering an effective amount of a compound of the formula:

or pharmaceutically acceptable salt thereof, in combination or alternation with an effective amount of 9-[2-(phosphono-methoxy)ethyl]adenine (PMEA, adefovir, dipivoxil), or pharmaceutically acceptable salt thereof.





98.

(Once Amended) A method for the treatment of a hepatitis B virus infection in a human comprising administering an effective amount of a compound of the formula:

or pharmaceutically acceptable salt thereof, in combination or alternation with an effective amount of lobucavir, or pharmaceutically acceptable salt thereof.

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(Once Amended) A method for the treatment of a hepatitis B virus infection in a human comprising administering an effective amount of a compound of the formula:

or pharmaceutically acceptable salt thereof, in combination or alternation with an effective amount of ganciclovir, or pharmaceutically acceptable salt thereof.





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(Once Amended) A method for the treatment of a hepatitis B virus infection in a human comprising administering an effective amount of a compound of the formula:

or pharmaceutically acceptable salt thereof, in combination or alternation with an effective amount of ribavirin, or pharmaceutically acceptable salt thereof.

(Once Amended) A method for the treatment of a hepatitis B virus infection in a human comprising administering an effective amount of a compound of the formula:

or pharmaceutically acceptable salt thereof, in combination or alternation with an effective amount of β -L-2-hydroxymethyl-5-(cytosin-1-yl)-1,3-oxathiolane (3TC), or pharmaceutically acceptable salt thereof.





HO 32.

(Once Amended) A method for the treatment of a hepatitis B virus infection in a human comprising administering an effective amount of a compound of the formula:

or pharmaceutically acceptable salt thereof, in combination or alternation with an effective amount of *cis*-2-hydroxymethyl-5-(5-fluorocytosin-1-yl)-1,3-oxathiolane (FTC), or pharmaceutically acceptable salt thereof.

53. (C hu

(Once Amended) A method for the treatment of a hepatitis B virus infection in a human comprising administering an effective amount of a compound of the formula:

or pharmaceutically acceptable salt thereof, in combination or alternation with an effective amount of β -L-2'-fluoro-5-methyl-arabinofuranosyl-uridine (L-FMAU), or pharmaceutically acceptable salt thereof.





(Once Amended) A method for the treatment of a hepatitis B virus infection in a human comprising administering an effective amount of a compound of the formula:

or pharmaceutically acceptable salt thereof, in combination or alternation with an effective amount of β -D-2,6-diaminopurine dioxolane (DAPD), or pharmaceutically acceptable salt thereof.

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(Once Amended) A method for the treatment of a hepatitis B virus infection in a human comprising administering an effective amount of a compound of the formula:

or pharmaceutically acceptable salt thereof, in combination or alternation with an effective amount of famciclovir, or pharmaceutically acceptable salt thereof.



(Once Amended) A method for the treatment of a hepatitis B virus infection in a human comprising administering an effective amount of a compound of the formula:

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or pharmaceutically acceptable salt thereof, in combination or alternation with an effective amount of penciclovir, or pharmaceutically acceptable salt thereof.

(Once Amended) A method for the treatment of a hepatitis B virus infection in a human comprising administering an effective amount of a compound of the formula:

or pharmaceutically acceptable salt thereof, in combination or alternation with an effective amount of 2-amino-1,9-dihydro-9-[4-hydroxy-3-(hydroxymethyl)-2-methylene-cyclopentyl]-6H-purin-6-one (entecavir, BMS-200475), or pharmaceutically acceptable salt thereof.





(Once Amended) A method for the treatment of a hepatitis B virus infection in a human comprising administering an effective amount of a compound of the formula:

or pharmaceutically acceptable salt thereof, in combination or alternation with an effective amount of 9-[2-(phosphono-methoxy)ethyl]adenine (PMEA, adefovir, dipivoxil), or pharmaceutically acceptable salt thereof.

(Once Amended) A method for the treatment of a hepatitis B virus infection in a human comprising administering an effective amount of a compound of the formula:

or pharmaceutically acceptable salt thereof, in combination or alternation with an effective amount of lobucavir, or pharmaceutically acceptable salt thereof.





(Once Amended) A method for the treatment of a hepatitis B virus infection in a human comprising administering an effective amount of a compound of the formula:

or pharmaceutically acceptable salt thereof, in combination or alternation with an effective amount of ganciclovir, or pharmaceutically acceptable salt thereof.

(Once Amended) A method for the treatment of a hepatitis B virus infection in a human comprising administering an effective amount of a compound of the formula:

or pharmaceutically acceptable salt thereof, in combination or alternation with an effective amount of ribavirin, or pharmaceutically acceptable salt thereof.

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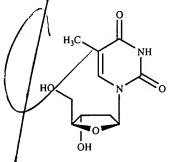




63. (New) A method for the prophylaxis of a hepatitis B virus infection in a human comprising administering an effective amount of β -L-2'-deoxycytidine of the formula:

or pharmaceutically acceptable salt thereof.

64. (New) A method for the prophylaxis of a hepatitis B virus infection in a human comprising administering an effective amount of β-L-thymidine of the formula:



or pharmaceutically acceptable salt thereof.

65. (New) A method for the prophylaxis of a hepatitis B virus infection in a human comprising administering an effective amount of a combination of the following nucleosides:

or a pharmaceutically acceptable salt thereof.



66. (New) A method for the prophylaxis of a hepatitis B virus infection in a human comprising administering an effective amount of a compound of the formula:

or pharmaceutically acceptable salt thereof, in combination or alternation with an effective amount of a compound selected from the group consisting of β-L-2-hydroxymethyl-5-(cytosin-1-yl)-1,3-oxathiolane (3TC), *cis*-2-hydroxymethyl-5-(5-fluorocytosin-1-yl)-1,3-oxathiolane (FTC), β-L-2'-fluoro-5-methyl-arabinofuranosyl-uridine (L-FMAU), β-D-2,6-diaminopurine dioxolane (DAPD), famciclovir, penciclovir, 2-amino-1,9-dihydro-9-[4-hydroxy-3-(hydroxymethyl)-2-methylenecyclopentyl]-6H-purin-6-one (entecavir BM\$-200475), 9-[2-(phosphono-methoxy)ethyl]adenine (PMEA, adefovir, dipivoxil); lobucavir, ganciclovir and ribavirin.

67. (New) A method for the prophylaxis of a hepatitis B virus infection in a human comprising administering an effective amount of a compound of the formula:

or pharmaceutically acceptable salt thereof, in combination or alternation with an effective amount of a compound selected from the group consisting of β -L-2-hydroxymethyl-5-(cytosin-1-yl)-1,3-oxathiolane (3TC), cis-2-hydroxymethyl-5-(5-fluorocytosin-1-yl)-1,3-oxathiolane (FTC), β -L-2'-fluoro-5-methyl-arabinofuranosyl-uridine (L-FMAU), β -D-2,6-diaminopurine dioxolane (DAPD), famciclovir, penciclovir,

